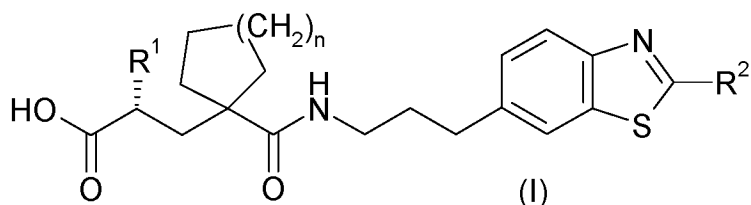


Claims:

1. (Original) A compound of formula (I)



wherein:

R¹ is H or CH₃;

R² is C₁-C₂ alkyl; and

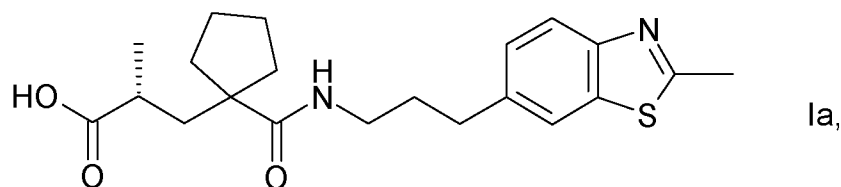
n is 1 or 2;

a tautomer thereof or a pharmaceutically acceptable salt, or solvate of said compound or tautomer.

2. (Original) The compound according to Claim 1 wherein n is 1.
3. (Original) The compound according to Claim 1 or Claim 2 wherein R¹ is hydrogen.
4. (Original) The compound according to Claim 3 wherein R² is methyl.
5. (Original) The compound according to Claim 3 wherein R² is ethyl.
6. (Original) The compound according Claim 1 or Claim 2 wherein R¹ is methyl.
7. (Original) The compound according to Claim 6 wherein R² is methyl.
8. (Original) The compound according to Claim 6 wherein R² is ethyl.
9. (Original) The compound according to Claims 1 or 2 wherein R² is methyl.
10. (Original) The compound according to Claims 1 or 2 wherein R² is ethyl.
11. (Original) The compound according to Claim 1 selected from
(*R*)-2-Methyl-3-(1-[[3-(2-methyl-1,3-benzothiazol-6-yl)propyl]carbamoyl]
cyclopentyl)propanoic acid;
3-(1-[[3-(2-ethyl-1,3-benzothiazol-6-yl)propyl]carbamoyl]cyclopentyl)propanoic
acid;
(*R*)-2-Methyl-3-(1-[[3-(2-ethyl-1,3-benzothiazol-6-yl)propyl]carbamoyl]
cyclopentyl)propanoic acid; or

3-(1-[[3-(2-ethyl-1,3-benzothiazol-6-yl)propyl]carbamoyl]cyclohexyl)propanoic acid.

12. (Original) A compound of Formula Ia,



a tautomer thereof or a pharmaceutically acceptable salt, or solvate of said compound or tautomer.

13. (Original) A pharmaceutical composition comprising a compound of formula (I) as claimed in any one of Claims 1, 2, 11 or 12, or pharmaceutically acceptable salts or solvates thereof, and a pharmaceutically acceptable diluent or carrier.

14. (Cancelled).

15. (Currently amended) A method of treating or preventing a disorder or condition ~~by inhibiting NEP~~ selected from obesity, female sexual dysfunction (FSD), sexual arousal disorder, female sexual arousal disorder (FSAD), male sexual dysfunction (MSD), male erectile dysfunction (MED), hypoactive sexual desire disorder, orgasmic disorder and sexual pain disorder in a mammal, comprising administering to said mammal a therapeutically effective amount of a compound of formula (I) as claimed in any one of Claims 1, 2, 11 or 12, or a pharmaceutically acceptable salt, or solvate thereof.

16. (Original) The method according to Claim 15 wherein n is 1.

17. (Original) The method according to Claim 15 or Claim 16 wherein R¹ is hydrogen.

18. (Original) The method according to Claim 17 wherein R² is methyl.

19. (Original) The method according to Claim 17 wherein R² is ethyl.

20. (Original) The method according to Claims 15 or 16 wherein R¹ is methyl.

21. (Original) The method according to Claim 20 wherein R² is methyl.

22. (Original) The method according to Claim 22 wherein R² is ethyl.

23. (Original) The method according to Claims 15 or 16 wherein R² is methyl.

24. (Original) The method according to Claims 15 or 16 wherein R² is ethyl.

25. (Cancelled).

26. (Currently Amended) The method according to Claim 15 25 wherein the disorder or condition is selected from female sexual dysfunction (FSD), sexual arousal disorder, female sexual arousal disorder (FSAD), male sexual dysfunction (MSD), male

erectile dysfunction (MED), hypoactive sexual desire disorder, orgasmic disorder and sexual pain disorder.

27. (Original) The method according to Claim 26 wherein the disorder or condition is selected from female sexual dysfunction (FSD), female sexual arousal disorder (FSAD), male sexual dysfunction (MSD), and male erectile dysfunction (MED).

28-33. (Cancelled).